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	7		10	Arai T. et al., "New antibiotics saframycins A, B, C, D and E," J Antibiot. (Tokyo) 1977, Vol. 30, No. 11, p.p. 1015-1018;												
	8			Bobbitt, J. et al., "Isoquinolines. III. A New Synthesis of 1,2,3,4-tetrahydro isoquinolines," J. Org. Chem. 1965, Vol. 30, p.p. 2247-2250;												
	9			Cabre-Castellvi, J. et al., "Convenient Synthesis of Carboxilic Acid Anhydrides using N,N-Bis[2-oxo-3-oxazol idinyl]phosphorodiamidic Chloride," <i>Synthesis</i> 1981 , No. 7, p.p. 616-620;												
	10		Caldwell C. et al., "Synthesis of the Lipophilic Side Chain of the Cyclic Hexa-depsipeptide Antibiotic L-156, 602," J. Org. Chem. 1990, Vol. 44, p.p. 2355-2361;													
	11		Caron, M. et al., "Highly Enantioselective Solvolyses of L- and D-Phenylalanine p-Nitrophenyl Esters by an L-Histidyl Dipeptide in Surfacant Coaggregates Formed by Cholesterol-Containing Amphiphiles," J. Org. Chem. 1988, Vol. 53, No. 21, p.p. 5187-5189;													
	12		Corey, E. et al., "Enantioselective Total Synthesis of Ecteinacidin 743," J. Am. Chem. Soc. 1996, Vol. 118, p.p 9202-9203;													
	13		Danishefsky, S. et al., "Total synthesis of Quinocarcinol Methyl Ester," J. Am. Chem. Soc. 1985, Vol. 107, No. 5, p.p. 1421-1423;													
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14	\	Fukuyama, T. et al., "Total Synthesis of(±) Saframycin A," J. Am. Chem. Soc. 1990, Vol. 112, No. 8, p.p. 3712-3713;												
15		Fukuyama, T. et al., "A Sterocontrolled Total Synthesis of (±) Reniramycin A," <i>Tetrahedron Lett.</i> 1990, Vol. 31 No. 42, p.p. 5989-5992;												
16		Fukuyama, T. et al., "Stereocontrolled total Synthesis of (±) Saframycin B," J. Am. Chem. Soc. 1982, Vol. 104 No. 118, p.p. 4957-4958;												
17	7	Gao, Y. et al., "Catalytic Asymmetric Epoxidation and Kinetic Resolution: Modified Procedures Including in Situ Derivatization," J. Am. Chem. Soc. 1987, Vol. 109, No. 18, p.p. 5765-5780;												
18		Guan, Y. et al., "Molecular and crystal structures of ecteinascidins: potent antitumor compounds from the Caribbean tunicate Ecteinascidia tur binata," <i>J. Biomol Struct. Dyn.</i> 1993, Vol. 10, No. 5, p.p. 793-817;												
19		Kishi, K. et al., "Structure-activity relationships of saframycins," J Antibiot. (Tokyo) 1984, Vol. 37, No. 8, p.p 847-852;												
20		Kitahara, Y. et al., "Synthesis of 4,7-Indolequinones. The Oxidative Demethylation of 4,7 Dimethoxyindoles with Ceric Ammonium Nitrate," <i>Chem. Phar. Bull. (Japan)</i> 1985, Vol. 33, No. 5, p.p. 2122-2128;												
21	l	Kubo, A. et al., "Stereoselective total Synthesis of (±) Saframycin B," J. Org. Chem. 1988, Vol. 53, No. 18, p.p. 4295-4310;												
22	2	Martinez, E. et al., "Phthalascidin, a synthetic antitumor agent with potency and mode of action comparable to ecteinacidin 743," <i>Proc. Natl. Acad. Sci.</i> 1999, Vol. 96, p.p. 3496-3501;												
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Serial No. Atty. Docket No. Form PTO-1449 **U.S. Department of Commerce** Not Yet Known 61071-AZ/ Patent and Trademark Office JPW/GJG/ACK **Applicants** INFORMATION DISCLOSURE CITATION Samuel J. Danishefsky et al. (Use several sheets if necessary) Filing Date Group 1624 Herewith U.S. PATENT DOCUMENTS Filing Date if Examiner Date Class Subclass **Document Number** Name Appropriate Initial / Item No. FOREIGN PATENT DOCUMENTS **Translation Document Number** Date Country Class Subclass Yes No OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.) Medina, E. et al., "Enantioselective synthesis of N-Boc-1-naphthylglycine," Tetrahedron Asym. 1997, Vol. 8, No. 23 10, p.p. 1581-1586; Mikami, Y. et al., "Saframycin S, a new saframycin group antibiotic," J. Pharmacobiodyn. 1981, No. 4, p.p. 282-24 Myers, A. et al. "A concise, Stereocontrolled Synthesis of (-) Saframycin A by the Directed Condensation of a-25 Amino Aldehyde Precursors," J. Am. Chem. Soc. 1999, Vol. 121, No. 43, p.p. 10828-10829; Sakai, R. et al., "Additional antitumor ecteinacidins from a Caribbean tunicate: Crystal structures and activities in 26 vivo," Proc. Natl. Acad. Sci. 1992, Vol. 89, p.p. 11456-11460; Sakai, R. et al., "Ecteinascidins: Putative Biosynthetic Precursors and Absolute Stereochemistry," J. Am. Chem. 27 Soc. 1996, Vol. 118, No. 35, p.p. 9017-9023; Sharpless, K. B. et al., "The Osmium-Catalyzed Asymmetric Dihydroxylation: A New Ligand Class and a Process 28 Improvement," J. Org. Chem. 1992, Vol., 57, No. 6, p.p. 2768-2771; Zhou et al., "A novel face specific Mannich closure providing access to the saframycin-ecteinascidin series of 29 piperazine based alkaloids," Tetrahedron Letters 2000, Vol. 41, p.p. 2043-2046; Zhou et al., "Synthetic explorations in the saframycin ecteinascidin series: construction of major chiral subunits 30 through catalytic asymmetric induction," Tetrahedron Letters 2000, Vol. 41, p.p. 2039-2042. EXAMINER DATE CONSIDERED

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